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      2
                  CA/CAplus records now contain indexing from 1907 to the
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      3
         SEP 09
                  present
         Jul 15
                  Data from 1960-1976 added to RDISCLOSURE
NEWS
                  Identification of STN records implemented
NEWS
         Jul 21
                  Polymer class term count added to REGISTRY
NEWS
         Jul 21
                  INPADOC: Basic index (/BI) enhanced; Simultaneous Left and
NEWS
      7
         Jul 22
                  Right Truncation available
                  New pricing for EUROPATFULL and PCTFULL effective
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      8
         AUG 05
                  August 1, 2003
                  Field Availability (/FA) field enhanced in BEILSTEIN
NEWS
      9
         AUG 13
                  PATDPAFULL: one FREE connect hour, per account, in
         AUG 15
NEWS 10
                  September 2003
                  PCTGEN: one FREE connect hour, per account, in
NEWS 11
         AUG 15
                  September 2003
                  RDISCLOSURE: one FREE connect hour, per account, in
         AUG 15
NEWS 12
                  September 2003
                  TEMA: one FREE connect hour, per account, in
NEWS 13
         AUG 15
                  September 2003
                  Data available for download as a PDF in RDISCLOSURE
NEWS 14
         AUG 18
NEWS 15
         AUG 18
                  Simultaneous left and right truncation added to PASCAL
                  FROSTI and KOSMET enhanced with Simultaneous Left and Righ
NEWS 16
         AUG 18
                  Truncation
                  Simultaneous left and right truncation added to ANABSTR
NEWS 17
         AUG 18
                  DIPPR file reloaded
NEWS 18
         SEP 22
                  INPADOC: Legal Status data to be reloaded
NEWS 19
         SEP 25
                  DISSABS now available on STN
NEWS 20
         SEP 29
NEWS EXPRESS April 4 CURRENT WINDOWS VERSION IS V6.01a, CURRENT
              MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
              AND CURRENT DISCOVER FILE IS DATED 01 APRIL 2003
              STN Operating Hours Plus Help Desk Availability
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FILE 'HOME' ENTERED AT 14:44:56 ON 30 SEP 2003

=> file medline, uspatful, dgene, wpids, fsta, biosis
COST IN U.S. DOLLARS
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0.21

FILE 'MEDLINE' ENTERED AT 14:45:19 ON 30 SEP 2003

FILE 'USPATFULL' ENTERED AT 14:45:19 ON 30 SEP 2003 CA INDEXING COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'DGENE' ENTERED AT 14:45:19 ON 30 SEP 2003 COPYRIGHT (C) 2003 THOMSON DERWENT

FILE 'WPIDS' ENTERED AT 14:45:19 ON 30 SEP 2003 COPYRIGHT (C) 2003 THOMSON DERWENT

FILE 'FSTA' ENTERED AT 14:45:19 ON 30 SEP 2003 COPYRIGHT (C) 2003 International Food Information Service

FILE 'BIOSIS' ENTERED AT 14:45:19 ON 30 SEP 2003 COPYRIGHT (C) 2003 BIOLOGICAL ABSTRACTS INC.(R)

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                     SHIMKETS RICHARD AUGUST/AU
                     SHIMKETS RICK/AU
E2
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E4
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Е6
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AB

L1 ANSWER 1 OF 4 USPATFULL on STN

Atrial natriuretic factor mutants and ischemic stroke
The present invention is based upon the observation that a mutant atrial
natriuretic factor (ANF) gene increases stroke latency in spontaneously
hypertensive rats-stroke prone (SHRSP). Accordingly, the present
invention provides methods using mutant ANF proteins, fragments,
analogs, derivatives and homologs of mutant ANF proteins, the nucleic
acids encoding these mutant ANF proteins, as well as modulators of ANF
for treating or preventing ischemic diseases, in particular ischemic
stroke. The invention also relates to methods of diagnosis, prognosis
and screening for a disposition for diseases and disorders associated
with increased levels of ANF. Pharmaceutical compositions, methods of
screening for ANF mutants and ANF modulators with utility for treatment
and prevention of ischemic stroke are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT. ACCESSION NUMBER: 2003:33456 USPATFULL

TITLE: Atrial natriuretic factor mutants and ischemic stroke

Shimkets, Richard August, West Haven, CT,

United States

CuraGen Corporation, New Haven, CT, United States (U.S. PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE B1 20030204 US 6514939

PATENT INFORMATION: APPLICATION INFO.: US 1999-428929 19991028 (9)

Continuation of Ser. No. US 1997-916043, filed on 21 RELATED APPLN. INFO.:

Aug 1997, now patented, Pat. No. US 6013630

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

Allen, Marianne P. PRIMARY EXAMINER:

LEGAL REPRESENTATIVE: Mintz Levin, Biswas, Naomi S., Elrifi, Ivor R.

18 NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

INVENTOR(S):

13 Drawing Figure(s); 10 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT: 2353

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 2 OF 4 USPATFULL on STN L1

TIAtrial natriuretic factor mutants and ischemic stroke

AΒ The present invention is based upon the observation that a mutant atrial natriuretic factor (ANF) gene increases stroke latency in spontaneously hypertensive rats-stroke prone (SHRSP). Accordingly, the present invention provides methods using mutant ANF proteins, fragments, analogs, derivatives and homologs of mutant ANF proteins, the nucleic acids encoding these mutant ANF proteins, as well as modulators of ANF for treating or preventing ischemic diseases, in particular ischemic stroke. The invention also relates to methods of diagnosis, prognosis and screening for a disposition for diseases and disorders associated with increased levels of ANF. Pharmaceutical compositions, methods of screening for ANF mutants and ANF modulators with utility for treatment and prevention of ischemic stroke are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT. ACCESSION NUMBER: 2000:4792 USPATFULL

Atrial natriuretic factor mutants and ischemic stroke TITLE:

INVENTOR (S): Shimkets, Richard August, West Haven, CT,

United States

PATENT ASSIGNEE(S): CuraGen Corporation, New Haven, CT, United States (U.S.

corporation)

NUMBER KIND DATE PATENT INFORMATION:

US 6013630 20000111 US 1997-916043 19970821 (8) APPLICATION INFO.:

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Allen, Marianne P.

LEGAL REPRESENTATIVE: Elrifi, Ivor R.Mintz, Levin, Cohn, Ferris, Glovsky and

Popeo P.C., Johnson, David E.

NUMBER OF CLAIMS: 2.8 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 12 Drawing Figure(s); 10 Drawing Page(s)

LINE COUNT: 2390

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 3 OF 4 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN

Atrial natriuretic factor mutants and ischemic stroke. TI

The present invention is based upon the observation that a mutant atrial AB natriuretic factor (ANF) gene increases stroke latency in spontaneously

hypertensive rats-stroke prone (SHRSP). Accordingly, the present invention provides methods using mutant ANF proteins, fragments, analogs, derivatives and homologs of mutant ANF proteins, the nucleic acids encoding these mutant ANF proteins, as well as modulators of ANF for treating or preventing ischemic diseases, in particular ischemic stroke. The invention also relates to methods of diagnosis, prognosis and screening for a disposition for diseases and disorders associated with increased levels of ANF. Pharmaceutical compositions, methods of screening for ANF mutants and ANF modulators with utility for treatment and prevention of ischemic stroke are also provided.

ACCESSION NUMBER: DOCUMENT NUMBER:

2003:129778 BIOSIS PREV200300129778

TITLE:

Atrial natriuretic factor mutants and ischemic stroke.

AUTHOR(S):

Shimkets, Richard August

ASSIGNEE: CuraGen Corporation

PATENT INFORMATION: US 6514939 February 04, 2003

SOURCE:

Official Gazette of the United States Patent and Trademark Office Patents, (Feb. 4 2003) Vol. 1267, No. 1, pp. No Pagination. http://www.uspto.gov/web/menu/patdata.html.

e-file.

ISSN: 0098-1133.

DOCUMENT TYPE: LANGUAGE:

Patent English

ANSWER 4 OF 4 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN Ь1.

Atrial natriuretic factor mutants and ischemic stroke. TI

The present invention is based upon the observation that a mutant atrial natriuretic factor (ANF) gene increases stroke latency in spontaneously AΒ hypertensive rats-stroke prone (SHRSP). Accordingly, the present invention provides methods using mutant ANF proteins, fragments, analogs, derivatives and homologs of mutant ANF proteins, the nucleic acids encoding these mutant ANF proteins, as well as modulators of ANF for treating or preventing ischemic diseases, in particular ischemic stroke. The invention also relates to methods of diagnosis, prognosis and screening for a disposition for diseases and disorders associated with increased levels of ANF. Pharmaceutical compositions, methods of screening for ANF mutants and ANF modulators with utility for treatment and prevention of ischemic stroke are also provided.

ACCESSION NUMBER:

2000:319969 BIOSIS

DOCUMENT NUMBER:

PREV200000319969

TITLE:

Atrial natriuretic factor mutants and ischemic stroke.

AUTHOR (S):

Shimkets, Richard August (1)

CORPORATE SOURCE:

(1) West Haven, CT USA

ASSIGNEE: CuraGen Corporation, New Haven, CT, USA

SOURCE:

PATENT INFORMATION: US 6013630 January 11, 2000 Official Gazette of the United States Patent and Trademark

> Office Patents, (Jan. 11, 2000) Vol. 1230, No. 2, pp. No pagination. e-file.

ISSN: 0098-1133.

DOCUMENT TYPE:

LANGUAGE:

Patent English

=> d l2 ti abs ibib tot

ANSWER 1 OF 2 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN  $L_2$ Novel G-protein-coupled receptors: Structure and tissue expression TTanalysis.

The G-protein-coupled receptors (GPCRs) are cell surface molecules that AΒ transmit extracellular signals to intracellular messengers. The GPCR gene family is the largest known receptor family including an estimated 700-1000 odorant GPCRs. The ligands that bind GPCRs are small and can be easily mimicked by use of synthetic analogues making GPCRs ideal targets for the development of therapeutic agents. We have identified several

hundred GPCR-like genes in our effort to characterize novel drug targets. These genes undergo a systematic analysis. The genes are first cloned and then tissue expression profiles generated for both normal and disease conditions. Extensive in silico analysis of these genes is conducted at both the nucleotide and amino acid level to determine the homology and secondary structure. The tissue expression profiles are correlated with the DNA sequence for each gene to enable us to distinguish the non-expressed pseudogenes. Additional analysis is directed towards the study of the upstream DNA sequence from multiple GPCRs with similar tissue-expression profiles. This sequence is then evaluated for conserved residues/motifs/domains which may be associated with potential regulatory and promoter regions that define tissue-specific expression profiles. Results from these analyses will be presented.

2001:514720 BIOSIS ACCESSION NUMBER: DOCUMENT NUMBER: PREV200100514720

Novel G-protein-coupled receptors: Structure and tissue TITLE:

expression analysis.

Padigaru, Muralidhara (1); Taupier, Ray (1); Zerhusen, Bryan (1); Herrmann, John (1); Burgess, Catherine (1); Braverman, Mike (1); Leach, Martin (1); Shenoy, Suresh (1); AUTHOR(S):

Shimkets, Rick (1)

(1) CuraGen Corporation, Branford, CT USA CORPORATE SOURCE:

International Genome Sequencing and Analysis Conference, SOURCE:

(2000) Vol. 12, pp. 82. print.

Meeting Info.: 12th International Genome Sequencing and Analysis Conference Miami Beach, Florida, USA September

12-15, 2000

DOCUMENT TYPE: Conference

English LANGUAGE: SUMMARY LANGUAGE: English

ANSWER 2 OF 2 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN L2

Prioritization of targets by in silico profile generation. TI

Genes differentially expressed between normal and diseased states offer an AΒ attractive target for drug-based intervention. CuraGen's proprietary gene expression technologies (SeqCalling, GeneCalling, Pharmacogenomics) employ a sophisticated bioinformatics platform that leads to the rapid identification of potential human disease targets and toxicology markers. The GeneCalling and Pharmacogenomics processes compare the gene expression of defined animal, tissue or cellular models of disease or drug-response to appropriate controls. For both of these methods, identification of differentially expressed genes does not require prior knowledge of gene sequence, a direct contrast to conventional chip-based technologies. Downstream sequence analysis of these potential targets is supplemented by cutting-edge genomics technologies and our proprietary human sequence database consisting of more than 4 million sequences. Target selection is further refined through the integration of the target's gene expression profile with our extensive protein: protein interaction database and high-throughput tissue distribution characterization. A completed profile of the selected target can therefore be rapidly and inversely engineered from the gene expression data. This dramatically reduces the non-productive consumption of resources while increasing the efficiency of selection. Further validation by wet-lab analysis is thereby directed toward targets that have the highest potential for downstream drug development or therapeutic intervention.

ACCESSION NUMBER:

2001:494515 BIOSIS

DOCUMENT NUMBER:

PREV200100494515

TITLE: AUTHOR(S):

Prioritization of targets by in silico profile generation. Burgess, Catherine (1); Braverman, Mike (1); Leach, Martin

(1); Gould-Rothberg, Bonnie (1); Shimkets, Rick (1)

CORPORATE SOURCE:

(1) CuraGen Corporation, Branford, CT USA

SOURCE:

International Genome Sequencing and Analysis Conference,

(2000) Vol. 12, pp. 52. print.

Meeting Info.: 12th International Genome Sequencing and

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Analysis Conference Miami Beach, Florida, USA September
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12-15, 2000

DOCUMENT TYPE: Conference LANGUAGE: English English SUMMARY LANGUAGE:

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(FILE 'HOME' ENTERED AT 14:44:56 ON 30 SEP 2003)

FILE 'MEDLINE, USPATFULL, DGENE, WPIDS, FSTA, BIOSIS' ENTERED AT 14:45:19 ON 30 SEP 2003

E SHIMKETS, R/AU

L1 4 S E1 2 S E2 T.2

=> e fernandes, e/au

E11 FERNANDES Z Z/AU

FERNANDES ZSCHABER R M/AU E2 5

0 --> FERNANDES, E/AU E3

FERNANDESALNEMRI T/AU E4 10 E5 FERNANDESHJDE B/AU

E6 1 FERNANDESM J M/AU FERNANDEX A/AU F:7 1

1 FERNANDEX C/AU E8

FERNANDEX DE LA VEGA CARLOS SANCHEZ/AU E9 1

E10 1 FERNANDEX DEL PALACIO M J/AU

FERNANDEX E GARCIA/AU E11 1. 7 FERNANDEX FERNANDEZ M/AU E12

=> s PROX polypeptide

2 PROX POLYPEPTIDE

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ANSWER 1 OF 2 USPATFULL on STN L3

Novel amino acid sequences for human fetal brain-like polypeptides TT AB This application is drawn to novel amino acid sequences for mammalian polypeptides that have sequence similarity to fetal brain tissue protein. The polypeptides are novel secreted proteins 649 amino acids in length.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

2003:4266 USPATFULL ACCESSION NUMBER:

TITLE: Novel amino acid sequences for human fetal brain-like

polypeptides

Shimkets, Richard A., West Haven, CT, UNITED STATES INVENTOR(S):

Fernandes, Elma, Branford, CT, UNITED STATES

PATENT ASSIGNEE(S): CuraGen Corporation, New Haven, CT (U.S. corporation)

NUMBER KIND DATE \_\_\_\_\_\_ PATENT INFORMATION: A1 20030102 US 2003004310 A1 20011205 APPLICATION INFO .: US 2001-4551 (10)

Continuation of Ser. No. US 2000-635949, filed on 10 RELATED APPLN. INFO.:

Aug 2000, PENDING

NUMBER DATE

PRIORITY INFORMATION: US 1999-148433P 19990811 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: MINTZ, LEVIN, COHN, FERRIS,, GLOVSKY AND POPEO, P.C., One Financial Center, Boston, MA, 02111

NUMBER OF CLAIMS:

15 1

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

5 Drawing Page(s)

LINE COUNT:

6347

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 2 OF 2 WPIDS COPYRIGHT 2003 THOMSON DERWENT on STN Nucleic acids encoding secreted polypeptides, designated PROX

polypeptides, useful for treating a syndrome associated with a PROX-associated disorder, e.g. cancer.

AN 2001-147509 [15] WPIDS

AB WO 200110902 A UPAB: 20011129

NOVELTY - Isolated nucleic acids encoding secreted polypeptides, designated PROX polypeptides (i.e. a PRO polypeptide where X is an integer from 1 to 17), are new.

DETAILED DESCRIPTION - Isolated nucleic acids encoding secreted polypeptides, designated PROX polypeptides (i.e. a PRO polypeptide where X is an integer from 1 to 17), are new.

A nucleic acid (N1) encoding a PROX polypeptide is selected from:

(a) a nucleic acid encoding a mature form of a polypeptide selected from one of the 17 amino acid sequences (P1) defined in the specification;

- (b) a nucleic acid encoding a variant of a mature form of a sequence selected from P1, where one or more amino acid residues differ from the sequence of the mature form, provided that the variant differs in no more than 15% of the residues from the sequence of the mature form;
  - (c) a nucleic acid encoding an amino acid sequence selected from P1;
- (d) a nucleic acid encoding a variant of a sequence selected from P1, where one or more amino acid residues differ from the sequence of the mature form, provided that the variant differs in no more than 15% of the residues from the amino acid sequence selected from P1;
- (e) a nucleic acid fragment encoding at least a portion of an amino acid sequence selected from P1, or a variant of the polypeptide, where one or more residues in the variant differs from the sequence of the mature form, provided that the variant differs in no more than 15% of the residues from the amino acid sequence selected from P1; or
  - (f) a nucleic acid comprising the complement of (a)-(e). INDEPENDENT CLAIMS are also included for the following:
- (1) an isolated polypeptide (P2) comprising an amino acid sequence selected from:
  - (a) a mature form of a polypeptide selected from P1;
- (b) a variant of a mature form of a sequence selected from P1, where one or more amino acid residues differ from the sequence of the mature form, provided that the variant differs in no more than 15% of the residues from the sequence of the mature form;
  - (c) an amino acid sequence selected from P1;
- (d) a variant of a sequence selected from P1, where one or more amino acid residues differ from the sequence of the mature form, provided that the variant differs in no more than 15% of the residues from the amino acid sequence selected from P1;
  - (2) a vector comprising N1 operably linked to a promoter;
  - (3) a cell comprising the vector of (2);
  - (4) an antibody (Ab1) that immunospecifically binds to P2;
- (5) a method of determining the presence or amount of P2 in a sample, comprising:
  - (a) contacting the sample with Ab1; and
- (b) determining the presence or amount of Ab1 bound to the polypeptide, therefore determining the presence or amount of P2 in the sample;
- (6) a method of determining the presence or amount of N1 in a sample, comprising:
  - (a) contacting the sample with a probe that binds to N1; and
  - (b) determining the presence or amount of the probe bound to the

nucleic acid;

- (7) a method of identifying an agent that binds to P2, comprising contacting the polypeptide with the agent and determining whether the agent binds to the polypeptide;
- (8) a method of identifying an agent that modulates the expression or activity of P2, comprising:
  - (a) providing a cell expressing the polypeptide;
  - (b) contacting the cell with the agent; and
- (c) determining whether the agent modulates expression or activity of the polypeptide, where an alteration in expression or activity of the polypeptide indicates that the agent modulates expression or activity of the polypeptide;
- (9) a method for modulating the activity of P2, comprising contacting a cell sample expressing P2 with a compound that binds to the polypeptide in an amount sufficient to modulate the activity if the polypeptide;
- (10) a method of treating or preventing a PROX-associated disorder in a human, comprising administering P2, N1 or Ab1;
  - (11) a kit comprising P2, N1 or Ab1;
- (12) a method (M1) for screening for a modulator of activity or of latency or predisposition to a PROX-associated disorder, comprising:
- (a) administering a test compound to a test animal at increased risk for a PROX-associated disorder, where the test animal recombinantly expresses P2;
- (b) measuring the activity of the polypeptide in the test animal after administering the test compound; and
- (c) comparing the activity of the protein in the test animal with the activity of the polypeptide in a control animal not administered the compound, where a change in the activity of the polypeptide in the test animal relative to the control animal indicates the test compound is a modulator of latency of or predisposition to a PROX-associated disorder;
- (13) a method for determining the presence of or predisposition to a disease associated with altered levels of P2 in a first mammalian subject, comprising:
- (a) measuring the level of expression of the polypeptide in a sample from the first mammalian subject; and
- (b) comparing the amount of the polypeptide in the sample of step (a) to the amount of polypeptide present in a control sample from a second mammalian subject known not to have, or not to be predisposed to the disease, where an alteration in the expression level of the polypeptide in the first subject as compared to the control sample indicates the presence of or predisposition to the disease;
- (14) a method for determining the presence of or predisposition to a disease associated with altered levels of N1 in a first mammalian subject;
- (15) a method of treating a pathological state in a mammal, comprising administering a polypeptide in an amount that is sufficient to alleviate the pathological state, where the polypeptide has a sequence at least 95% identical to a polypeptide comprising an amino acid sequence selected from P1, or its biologically active fragment; and
- (16) a method of treating a pathological state in a mammal, comprising administering Ab1 in an amount that is sufficient to alleviate the pathological state.

ACTIVITY - Cytostatic; Immunomodulatory; reproduction general. No biological data given.

MECHANISM OF ACTION - Gene therapy; PROX antagonist; PROX agonist. USE - The PROX polypeptide, nucleic acid and antibody are useful in the manufacture of a medicament for treating a syndrome associated with a PROX-associated disorder (claimed), e.g. a cell proliferation and/or differentiation disorder (e.g. cancer or immune associated disorders) and a gestational disease (e.g. pre-clampsia).

They are also used for screening for a modulator of activity or of latency or predisposition to a PROX-associated disorder.

Dwg.0/9

ACCESSION NUMBER: DOC. NO. CPI:

TITLE:

Nucleic acids encoding secreted polypeptides, designated

PROX polypeptides, useful for treating a syndrome

associated with a PROX-associated disorder, e.g. cancer.

DERWENT CLASS:

B04 D16

95

INVENTOR(S):

FERNANDES, E; SHIMKETS, R A

PATENT ASSIGNEE(S):

(CURA-N) CURAGEN CORP

COUNTRY COUNT:

PATENT INFORMATION:

PATENT NO KIND DATE WEEK LA PG

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WO 2001010902 A2 20010215 (200115)\* EN 166

RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ

NL OA PT SD SE SL SZ TZ UG ZW

W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CR CU CZ DE DK DM

DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC

LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD SE

SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW

AU 2000068997 A 20010305 (200130)

EP 1218406 A1 20020703 (200251) EN

R: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT LU LV MC MK NL PT

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US 2003004310 A1 20030102 (200305)

JP 2003508030 W 20030304 (200319)

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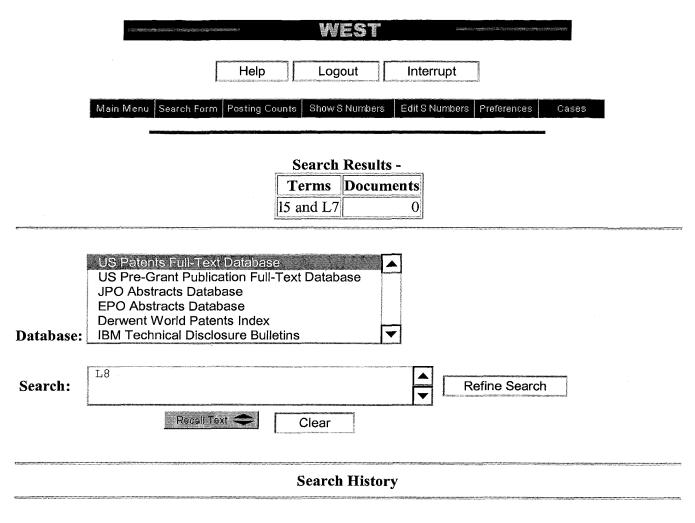
#### APPLICATION DETAILS:

PATENT NO KIND	AP:	PLICATION	DATE
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US 2003004310 A1 Pro Con	ovisional US	2000-US21857 1999-148433P 2000-635949	20000811 19990811 20000810
JP 2003508030 W	WO	2001-4551 2000-US21857 2001-515709	20011205 20000811 20000811

#### FILING DETAILS:

PATENT NO	KIND			PA.	TENT NO
AU 200006899	7 A	Based	on	WO.	2001010902
EP 1218406	A1	Based	on	WO	2001010902
JP 200350803	O W	Based	on	WO	2001010902

PRIORITY APPLN. INFO: US 2000-635949 20000810; US 1999-148433P 19990811; US 2001-4551 20011205



DATE: Tuesday, September 30, 2003 Printable Copy Create Case

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<u>L7</u>	11 and L6	15	<u>L7</u>
<u>L6</u>	fernandes.in.	193	<u>L6</u>
<u>L5</u>	11 and L4	5	<u>L5</u>
<u>L4</u>	Shimkets.in.	6	<u>L4</u>
<u>L3</u>	protease inhibitor and L2	36064	<u>L3</u>
<u>L2</u>	L1 and cancer	17499	<u>L2</u>
L1	PROX polypeptide	43127	L1

END OF SEARCH HISTORY

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## Search Results - Record(s) 1 through 10 of 15 returned.

1. Document ID: US 6566505 B2

L7: Entry 1 of 15

File: USPT

May 20, 2003

US-PAT-NO: 6566505

DOCUMENT-IDENTIFIER: US 6566505 B2

TITLE: Antibodies to Mch6 polypeptides

DATE-ISSUED: May 20, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA
Fernandes-Alnemri; Teresa Ambler PA
Litwack; Gerald Bryn Mawr PA

US-CL-CURRENT: 530/387.9; 530/388.26, 530/389.1

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | MMC | Draw, Desc | Image |

2. Document ID: US 6462175 B1

L7: Entry 2 of 15

File: USPT

Oct 8, 2002

US-PAT-NO: 6462175

DOCUMENT-IDENTIFIER: US 6462175 B1

TITLE: Mch3, a novel apoptotic protease, nucleic acids encoding and methods of use

DATE-ISSUED: October 8, 2002

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA
Fernandes-Alnemri; Teresa Ambler PA
Litwack; Gerald Wynnewood PA
Armstrong; Robert San Diego CA
Tomaselli; Kevin La Jolla CA

US-CL-CURRENT: 530/350; 435/226, 530/300

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMIC Draw Desc Image

3. Document ID: US 6455296 B2

L7: Entry 3 of 15

File: USPT

Sep 24, 2002

US-PAT-NO: 6455296

DOCUMENT-IDENTIFIER: US 6455296 B2

TITLE: Apoptotic protease Mch6, nucleic acids encoding same and methods of use

DATE-ISSUED: September 24, 2002

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA
Fernandes-Alnemri; Teresa Ambler PA
Litwack; Gerald Wynnewood PA

US-CL-CURRENT: 435/226; 435/219

#### Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWIC | Draw Desc | Image |

### 4. Document ID: US 6287795 B1

L7: Entry 4 of 15

File: USPT

Sep 11, 2001

US-PAT-NO: 6287795

DOCUMENT-IDENTIFIER: US 6287795 B1

TITLE: Mch4 and Mch5, apoptotic protease, nucleic acids encoding and methods of use

DATE-ISSUED: September 11, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA

Fernandes-Alnemri; Teresa Ambler PA

Litwack; Gerald Wynnewood PA

Armstrong; Robert San Diego CA

Tomaselli; Kevin La Jolla CA

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |

US-CL-CURRENT: 435/23; 435/219, 435/325, 435/7.21, 435/7.72, 435/7.91, 536/23.2

5. Document ID: US 6274318 B1

L7: Entry 5 of 15

File: USPT

Aug 14, 2001

KWIC Draw Deso Image

US-PAT-NO: 6274318

DOCUMENT-IDENTIFIER: US 6274318 B1

\*\* See image for Certificate of Correction \*\*

TITLE: Apoptotic protease Mch6, nucleic acids encoding same and methods of us

DATE-ISSUED: August 14, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA
Fernandes-Alnemri; Teresa Ambler PA
Litwack; Gerald Wynnewood PA

US-CL-CURRENT: 435/6; 435/226, 435/23, 435/7.6, 435/7.71, 435/7.72, 435/7.9

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | KMC | Draw. Desc | Image |

6. Document ID: US 6271361 B1

L7: Entry 6 of 15

File: USPT

Aug 7, 2001

US-PAT-NO: 6271361

DOCUMENT-IDENTIFIER: US 6271361 B1

TITLE: Apoptotic protease Mch6, nucleic acids encoding same and methods of use

DATE-ISSUED: August 7, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA
Fernandes-Alnemri; Teresa Ambler PA
Litwack; Gerald Wynnewood PA

US-CL-CURRENT: 536/23.2; 435/226, 435/320.1, 435/69.1, 536/23.5

Full Title Citation Front Review Classification Date Reference Sequences Attachments KMC Draw Desc Image

7. Document ID: US 5861498 A

L7: Entry 7 of 15

File: USPT

Jan 19, 1999

US-PAT-NO: 5861498

DOCUMENT-IDENTIFIER: US 5861498 A

\*\* See image for Certificate of Correction \*\*

TITLE: Nucleotides encoding immunophilin FKBP46 and fragments thereof

DATE-ISSUED: January 19, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA
Fernandes-Alnemri; Teresa Ambler PA
Litwack; Gerald Wynnewood PA

US-CL-CURRENT: 536/23.5; 435/320.1

Full Title Citation Front Review Classification Date Reference Sequences Attachments KMC Draw Desc Image

8. Document ID: US 5858778 A

L7: Entry 8 of 15

File: USPT

Jan 12, 1999

US-PAT-NO: 5858778

DOCUMENT-IDENTIFIER: US 5858778 A

TITLE: SF caspase-1 and compositions for making and methods of using the same

DATE-ISSUED: January 12, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA
Fernandes-Alnemri; Teresa Ambler PA
Litwack; Gerald Bryn Mawr PA

US-CL-CURRENT:  $\frac{435}{325}$ ;  $\frac{435}{219}$ ,  $\frac{435}{252.3}$ ,  $\frac{435}{252.33}$ ,  $\frac{435}{254.11}$ ,  $\frac{435}{320.1}$ ,  $\frac{435}{348}$ ,  $\frac{536}{23.2}$ ,  $\frac{536}{24.3}$ ,  $\frac{536}{24.31}$ 

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |

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9. Document ID: US 5851815 A

L7: Entry 9 of 15

File: USPT

Dec 22, 1998

US-PAT-NO: 5851815

DOCUMENT-IDENTIFIER: US 5851815 A

TITLE: MCH4 and MCH5, apoptotic proteases

DATE-ISSUED: December 22, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Alnemri; Emad S. Ambler PA

Fernandes-Alnemri; Teresa Ambler PA

Litwack; Gerald Wynnewood PA

Armstrong; Robert San Diego CA

Tomaselli; Kevin La Jolla CA

US-CL-CURRENT: 435/219; 435/183, 435/212, 530/324

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments

KMC Draw Desc Image

10. Document ID: US 5786173 A

L7: Entry 10 of 15

File: USPT

Jul 28, 1998

US-PAT-NO: 5786173

DOCUMENT-IDENTIFIER: US 5786173 A

TITLE: MCH4 and MCH5, apoptotic protease, nucleic acids encoding and methods of use

DATE-ISSUED: July 28, 1998

INVENTOR-INFORMATION:

NAME CITY ZIP CODE STATE COUNTRY Ambler Alnemri; Emad S. PΑ Fernandes-Alnemri; Teresa Ambler PΑ Litwack; Gerald Wynnewood PΑ Armstrong; Robert San Diego CA Tomaselli; Kevin La Jolla CA

Full	Title Citation Front	Review Classification Date F	eference Sequences	Attachments	KMMC   Drawn Desc   Image
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### Search Results - Record(s) 11 through 15 of 15 returned.

11. Document ID: US 5702699 A

L7: Entry 11 of 15

File: USPT

Dec 30, 1997

US-PAT-NO: 5702699

DOCUMENT-IDENTIFIER: US 5702699 A

TITLE: Process for the recovery of lipophilic proteins

DATE-ISSUED: December 30, 1997

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Hanisch; Wolfgang H.

Oakland

CA

Fernandes; Peter

Walnut Creek

CA

US-CL-CURRENT: 424/85.6; 530/351

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |

KWIC Draw Desc Image

☐ 12. Document ID: US 5643566 A

L7: Entry 12 of 15

File: USPT

Jul 1, 1997

US-PAT-NO: 5643566

DOCUMENT-IDENTIFIER: US 5643566 A

TITLE: Formulation processes for lipophilic proteins

DATE-ISSUED: July 1, 1997

INVENTOR-INFORMATION:

AME

CITY

STATE ZIP CODE

COUNTRY

Hanisch; Wolfgang H.

Balmoral Heights

AU

Fernandes; Pete M.

Walnut Creek

CA

Taforo; Terrance

Oakland

CA

Thomson; James W.

Albany

CA

US-CL-CURRENT: 424/85.4; 424/85.2, 424/85.6, 530/351

Full Title Citation Front Review Classification Date Reference Sequences Attachments

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☐ 13. Document ID: US 4992271 A

L7: Entry 13 of 15

File: USPT

Feb 12, 1991

US-PAT-NO: 4992271

DOCUMENT-IDENTIFIER: US 4992271 A

TITLE: Formulation for lipophilic IL-2 proteins

DATE-ISSUED: February 12, 1991

INVENTOR - INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Hanisch; Wolfgang H. Oakland CA
Fernandes; Peter M. Walnut Creek CA
Taforo; Terrance Oakland CA

US-CL-CURRENT: 424/85.2; 424/85.1, 424/85.4, 424/85.5, 424/85.6, 424/85.7, 435/811,

514/12, 514/2, 514/21, 514/8, 514/885, 514/970, 530/351

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments

☐ 14. Document ID: US 4569910 A

L7: Entry 14 of 15 File: USPT

Feb 11, 1986

KMC | Draw Desc | Image

US-PAT-NO: 4569910

DOCUMENT-IDENTIFIER: US 4569910 A

TITLE: Methods and reagents for pyranosone production

DATE-ISSUED: February 11, 1986

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Koths; Kirston E. Berkeley CA Halenbeck; Robert F. San Rafael CA Fernandes; Peter M. Walnut Creek CA

US-CL-CURRENT: 435/105; 435/190, 435/911

Full Title Citation Front Review Classification Date Reference Sequences Attachments KMC Draw Desc Image

15. Document ID: US 4462940 A

L7: Entry 15 of 15 File: USPT Jul 31, 1984

US-PAT-NO: 4462940

DOCUMENT-IDENTIFIER: US 4462940 A

TITLE: Process for the recovery of human .beta.-interferon-like polypeptides

DATE-ISSUED: July 31, 1984

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Hanisch; Wolfgang H. Oakland CA Fernandes; Peter M. Lafayette CA

US-CL-CURRENT: 530/351; 424/85.6, 435/69.51, 435/811, 530/417, 530/424, 530/808,

# 530/825

Full   Title   Citation   Front   Review   Classification   Date   Reference   Sequences   Attachments   KWC   Draw Desc   Image			
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## **Search Results -** Record(s) 1 through 5 of 5 returned.

1. Document ID: US 6610480 B1

L5: Entry 1 of 5

File: USPT

Aug 26, 2003

US-PAT-NO: 6610480

DOCUMENT-IDENTIFIER: US 6610480 B1

TITLE: Treatment and diagnosis of cardiac hypertrophy

DATE-ISSUED: August 26, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Shimkets; Richard A.

West Haven

CT

Lowe; David G.

Hillsborough

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw Desc Image

US-CL-CURRENT:  $\underline{435}/\underline{6}$ ;  $\underline{536}/\underline{23.1}$ ,  $\underline{536}/\underline{24.3}$ ,  $\underline{536}/\underline{24.31}$ ,  $\underline{536}/\underline{25.3}$ 

☐ 2. Document ID: US 6537554 B1

L5: Entry 2 of 5

File: USPT

Mar 25, 2003

US-PAT-NO: 6537554

DOCUMENT-IDENTIFIER: US 6537554 B1

TITLE: Nucleotide sequences and amino acid sequences of secreted proteins involved in angiogenesis

DATE-ISSUED: March 25, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Shimkets; Richard A.

West Haven

CT

Jeffers; Michael

Branford

CT

US-CL-CURRENT: 424/198.1; 424/184.1, 435/4, 436/64

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KNNC Draw. Desc Image

3. Document ID: US 6514939 B1

L5: Entry 3 of 5

File: USPT

Feb 4, 2003

US-PAT-NO: 6514939

DOCUMENT-IDENTIFIER: US 6514939 B1

TITLE: Atrial natriuretic factor mutants and ischemic stroke

DATE-ISSUED: February 4, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Shimkets; Richard August

West Haven

CT

US-CL-CURRENT: 514/12; 530/324, 530/350

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw Desc Image

4. Document ID: US 6486299 B1

L5: Entry 4 of 5

File: USPT

Nov 26, 2002

US-PAT-NO: 6486299

DOCUMENT-IDENTIFIER: US 6486299 B1

TITLE: Genes and proteins predictive and therapeutic for stroke, hypertension,

diabetes and obesity

DATE-ISSUED: November 26, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Shimkets; Richard A.

West Haven

CT

US-CL-CURRENT: 530/350; 530/380, 530/800

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KMC | Draw Desc | Image |

5. Document ID: US 6013630 A

L5: Entry 5 of 5

File: USPT

Jan 11, 2000

US-PAT-NO: 6013630

DOCUMENT-IDENTIFIER: US 6013630 A

TITLE: Atrial natriuretic factor mutants and ischemic stroke

DATE-ISSUED: January 11, 2000

INVENTOR - INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Shimkets; Richard August

West Haven

CT

US-CL-CURRENT: <u>514/12</u>; <u>530/324</u>

Full Title Citation Front Review Classification Date Reference Sequences Attachments

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